SUPPLEMENTAL DECLARATION removes the basis for the outstanding rejection.

Amendments to the claims are reflected in the listing of claims which begins on page 3 of this paper. They differ from the claims prior to the present amendment as follows:

The following pending claims are amended: 13, 28, 31 and 40.

The following claims which were pending prior to the present amendment are cancelled: 14, 15, 24, 29, 32, 33, 41, 43, 52, 61 and 79-86.

The REMARKS begin on page 28 of this paper.

Claim Amendment Directions:

Cancel claims 14, 15, 24, 29, 32, 33, 41, 43, 52, 61 and 79-86.

Amend claims 13, 28, 31 and 40 as set forth in the following listing of claims.

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Cancelled)

2. (Cancelled)

3. (Cancelled)

4. (Cancelled)

5. (Cancelled)

6. (Cancelled)

7. (Cancelled)

8. (Cancelled)

9. (Cancelled)

10. (Cancelled)

11. (Cancelled)

12. (Cancelled)

13. (Fourth Amendment) [The compound of claim 1, wherein:

R represents a hydrogen atom, a halogen atom or an alkyl group having from 1 to 4 carbon atoms;

R^1represents a methyl group, an amino group or an acetylamino group;

R^2represents an unsubstituted phenyl group or a phenyl group which is substituted by at least one substituent selected from the group consisting of a halogen atom: an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 4 carbon atoms; an alkyl group having from 1 to 4 carbon atoms which is substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthiogroup having from 1 to 4 carbon atoms; a mercapto group; an alkanoyl group having from 1 to 4 carbon atoms; a haloalkyl group having from 1 to 4 carbon atoms; and an alkylenedioxy group having from 1 to 4 carbon atoms;

R^3represents a hydrogen atom, a halogen atom, an unsubstituted alkyl group having from 1 to 4 carbon atoms or a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms;

R^4represents

a hydrogen atom;

an unsubstituted alkyl group having from 1 to 4 carbon atoms;

a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms;

a cycloalkyl group having from 3 to 6 carbon atoms;

an aryl group which has from 6 to 10 ring carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 6 carbon atoms; an alkyl group having from 1 to 6 carbon atoms and substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having 1 to 4 carbon atoms and an alkylthio group having 1 to 4 carbon atoms; and a cycloalkoxy group having 3 to 8 carbon atoms; and an aralkyl group having from 1 to 4 carbon atoms in the alkyl part and containing at least one said aryl group] A compound selected from the group consisting of the following eight compounds and a pharmaceutically acceptable salt of said compounds:

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole,

2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole, and

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole .

14. (Cancelled)

15. (Cancelled)

16. (First Amendment) [The compound of claim 1, which is 4-methyl] 4-Methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl) pyrrole.

17. (First Amendment) [The compound of claim 1, which is 2-(4-methoxyphenyl)] 2-(4-Methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

18. (First Amendment) [The compound of claim 1, which is 2-(4-chlorophenyl)] 2-(4-Chlorophenyl)-4-methyl-1-(4-

sulfamoylphenyl)pyrrole.

19. (First Amendment) [The compound of claim 1, which is 4-methyl] 4-Methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl) pyrrole.

20. (First Amendment) [The compound of claim 1, which is 2-(4-ethoxyphenyl)] 2-(4-Ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

21. (First Amendment) [The compound of claim 1, which is 2-(4-methoxy] 2-(4-Methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

22. (First Amendment) [The compound of claim 1, which is 2-(3-fluoro] 2-(3-Fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

23. (First Amendment) [The compound of claim 1, which is 2-(3,4-dimethylphenyl)] 2-(3,4-Dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

24. (Cancelled)

25. (Cancelled)

26. (Cancelled)

27. (Cancelled)

28. (Fourth Amendment) [The method of claim 27, wherein:

R represents a hydrogen atom, a halogen atom or an alkyl group having from 1 to 4 carbon atoms;

R^1 represents a methyl group, an amino group or an acetylamino group;

W represents

an unsubstituted phenyl group or;

a phenyl group which is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 4 carbon atoms; an alkyl group having from 1 to 4 carbon atoms and which is substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms; a mercapto group; an alkanoylthio group having from 1 to 4 carbon atoms; a haloalkoxy group having from 1 to 4 carbon atoms; and an alkylenedioxy group having from

1 to 4 carbon atoms;

R^3represents a hydrogen atom, a halogen atom, an unsubstituted alkyl group having from 1 to 4 carbon atoms or a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms;

R^4represents

a hydrogen atom;

an unsubstituted alkyl group having from 1 to 4 carbon atoms;

a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms;

a cycloalkyl group having from 3 to 6 carbon atoms;

an aryl group which has from 6 to 10 ring carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 3 carbon atoms; an alkyl group having from 1 to 3 carbon atoms and substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy

group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms; and a cycloalkyloxy group having from 3 to 8 carbon atoms; and

an aralkyl group having from 1 to 4 carbon atoms in the alkyl part and containing at least one said aryl group] A method of treating or relieving pain or inflammation in a mammal suffering therefrom comprising administering to a mammal in need thereof an effective anti-inflammatory amount or effective analgesic amount of a compound selected from the group consisting of the following eight compounds and a pharmaceutically acceptable salt of said compounds:

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl) pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole,

2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole, and

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole .

29. (Cancelled)

30. (Cancelled)

31. (Second Amendment) A method of inhibiting bone resorption in a mammal comprising administering to a mammal in need thereof a pharmaceutically effective amount of a compound selected from the group consisting of the following eight compounds [compound of formula (I), the compound of formula (II),] and a pharmaceutically acceptable salt of said compounds: [as claimed in claim 1]

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl) pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole,

2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole, and

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole .

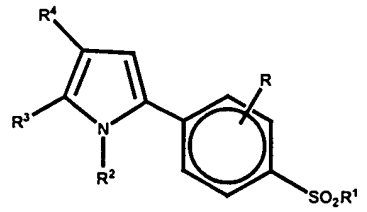
32. (Cancelled)

33. (Cancelled)

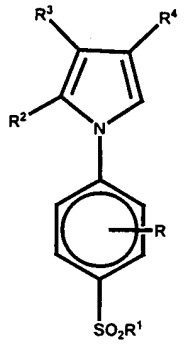
34. (Cancelled)

35. (First Amendment) A method of inhibiting leukotriene production in a mammal comprising administering to a mammal in need thereof a compound selected from the group consisting of the compound of formula (I), the compound of formula (II) and a pharmaceutically acceptable salt of said compound [as claimed in claim 1] wherein:

(I)



(II)



R represents a hydrogen atom, a halogen atom or an alkyl group having from 1 to 6 carbon atoms;

R^1 represents an alkyl group having from 1 to 6 carbon atoms or an amino group;

R^2 represents a phenyl group which is unsubstituted or is

substituted by at least one substituent selected from the group consisting of substituents α and subsituents β defined below;

R^3 represents a hydrogen atom, a halogen atom or an alkyl group which has from 1 to 6 carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms;

R^4 represents a hydrogen atom; an alkyl group which has from 1 to 6 carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms; a cycloalkyl group having from 3 to 8 carbon atoms, an aryl group which is as defined below, or an aralkyl group which is as defined below;

said aryl group having from 6 to 14 ring carbon atoms in a carbocyclic ring and are unsubstituted or are substituted by at least one substituent selected from the group consisting of substituents α and substituents β, defined below;

said aralkyl group are an alkyl group having from 1 to 6 carbon atoms and which are substituted by at least one aryl group as defined above;

said substitutents α are selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms; said substituents β are selected from the group consisting of an alkyl group which has from 1 to 6 carbon atoms and which is unsubstituted or are substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms; an alkanoyloxy group having from 1 to 6 carbon atoms; a mercapto group; an alkanoylthio group having from 1 to 6 carbon atoms; an alkylsulfinyl group having from 1 to 6 carbon atoms; a cycloalkyloxy group having from 3 to 8 carbon atoms; a haloalkoxy group having from 1 to 6 carbon atoms; and an alkylenedioxy group having from 1 to 6 carbon atoms;

or a pharmaceutically acceptable salt thereof.

36. (Third Amendment) The method of claim 35, wherein:

R represents a hydrogen atom, a halogen atom or an alkyl group having from 1 to 4 carbon atoms;

[R^2]

$\underline{R^1}$ represents a methyl group[,] <u>or</u> an amino group [or an acetylamino group];

R^2 represents

an unsubstituted phenyl group or a phenyl group which is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 4 carbon atoms; an alkyl group having from 1 to 4 carbon atoms and which is substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms; [a mercapto group; an alkanoylthio group having from 1 to 4 carbon atoms;] a haloalkoxy group having from 1 to 4 carbon atoms; and an alkylenedioxy group having from 1 to 4 carbon atoms;

R^3 represents a hydrogen atom, a halogen atom, an unsubstituted alkyl group having from 1 to 4 carbon

atoms or a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms;

R^4 represents

a hydrogen atom;

an unsubstituted alkyl group having from 1 to 4 carbon atoms;

a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to [6] 4 carbon atoms and an alkylthio group having from 1 to [6] 4 carbon atoms;

a cycloalkyl group having from 3 to 6 carbon atoms;

an aryl group which has from 6 to 10 ring carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to [6] 4 carbon atoms; an alkyl group having from 1 to [6] 4 carbon atoms and substituted by

at least one substituent selected from the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to [6] 4 carbon atoms and an alkylthio group having from 1 to [6] 4 carbon atoms; and a cycloalkyloxy group having from 3 to [8] 7 carbon atoms; an aralkyl group having from 1 to 4 carbon atoms in the alkyl part and containing at least one said aryl group.

37. (Third Amendment) The method of claim 35, wherein:

R represents a hydrogen atom[, a fluorine atom, a chlorine atom or a methyl group];

R^1 represents an amino group [or an acetylamino group];

R^2 represents

an unsubstituted phenyl group or

a phenyl group which is substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms, an alkylthio group having from 1 to 4 carbon atoms, an alkyl group having from 1 to 4 carbon atoms, a haloalkyl group having from 1 to 4 carbon atoms, [a mercapto group, an alkanoylthio group having from 1 to 4 carbon atoms,] a haloalkoxy group having from 1 to 4 carbon atoms and a alkylenedioxy group

having from 1 to 4 carbon atoms;

R^3 represents a hydrogen atom, a halogen atom, an alkyl group having from 1 to 4 carbon atoms or a haloalkyl group having from 1 to 4 carbon atoms;

R^4 represents

a hydrogen atom;

an unsubstituted alkyl group having from 1 to 4 carbon atoms;

a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a hydroxy group and an alkoxy group having from 1 to [6] 4 carbon atoms;

a cycloalkyl group having from 3 to 6 carbon atoms;

an aryl group which has from 6 to 10 ring carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of a hydroxy group; a halogen atom; an alkoxy group having from 1 to [6] 4 carbon atoms; an unsubstituted alkyl group having from 1 to [6] 4 carbon atoms; an alkyl group having from 1 to [6] 4 carbon atoms and which is unsubstituted or substituted by at least one halogen atom; and a cycloalkyloxy group having from 3 to [8] 7 carbon atoms; and an aralkyl group having from 1 to 4 carbon atoms in the

alkyl part and containing at least one said aryl group.

38. (Cancelled)

39. (Cancelled)

40. (Fourth Amendment) [The method of claim 39, wherein:

R represents a hydrogen atom, a halogen atom or an alkyl group having from 1 to 4 carbon atoms;

R^1represents a methyl group, an amino group or an acetylamino group;

R^2represents

an unsubstituted phenyl group or

a phenyl group which is substituted by at least one substituent selected from the group consisting of a halogen atom; an alkoxy group having 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 4 carbon atoms; an alkyl group having from 1 to 4 carbonatoms and which is substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms; a mercapto group; an alkanoylthio group having from 1 to 4 carbon atoms; a haloalkoxy group having from

1 to 4 carbon atoms; and an alkylenedioxy group having from 1 to 4 carbon atoms;

R^3represents a hydrogen atom, a halogen atom, an unsubstituted alkyl group having from 1 to 4 carbon atoms or a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected from the group consisting of a halogen atom, an alkoxy group having from 1 to 4 carbon atoms and an alkylthio group having from 1 to 4 carbon atoms;

R^4represents

a hydrogen atom;

an unsubstituted alkyl group having from 1 to 4 carbon atoms;

a substituted alkyl group having from 1 to 4 carbon atoms and substituted by at least one substituent selected fromthe group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthiogroup having from 1 to 6 carbon atoms;

a cycloalkyl group having from 3 to 6 carbon atoms;

an aryl group which has from 6 to 10 ring carbon atoms and which is unsubstituted or is substituted by at least one substituent selected from the group consisting of ahalogen atom; an alkoxy group having from 1 to 4 carbon atoms; an alkylthio group having from 1 to 4 carbon atoms; an unsubstituted alkyl group having from 1 to 6 carbon atoms; an alkyl group having from 1 to 6 carbon atoms and substituted by at least one substituent selected from

the group consisting of a hydroxy group, a halogen atom, an alkoxy group having from 1 to 6 carbon atoms and an alkylthio group having from 1 to 6 carbon atoms; and a cycloalkyloxy group having from 3 to 8 carbon atoms; and

an aralkyl group having from 1 to 4 carbon atoms in thealkyl part and containing at least one said aryl group] A method of selectively inhibiting the activity of COX-2 in a mammal comprising administering to said mammal a pharmaceutically effective amount of a compound selected from the group consisting of the following eight compounds and a pharmaceutically acceptable salt of said compounds:

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl) pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole,

2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole, and

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole .

41. (Cancelled)

42. (Cancelled)

43. (Cancelled)

44. (First Amendment) The method of claim 28 wherein said compound is 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl) pyrrole.

45. (First Amendment) The method of claim 28 wherein said compound is 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

46. (First Amendment) The method of claim 28 wherein said compound is 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

47. (First Amendment) The method of claim 28 wherein said compound is 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl) pyrrole.

48. (First Amendment) The method of claim 28 wherein said compound is 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)

pyrrole.

49. (First Amendment) The method of claim 28 wherein said compound is 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

50. (First Amendment) The method of claim 28 wherein said compound is 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

51. (First Amendment) The method of claim 28 wherein said compound is 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

52. (Canceled)

53. (New) The method of claim 31 wherein said compound is 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole.

54. (New) The method of claim 31 wherein said compound is 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

55. (New) The method of claim 31 wherein said compound is 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

56. (New) The method of claim 31 wherein said compound is 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole.

57. (New) The method of claim 31 wherein said compound is 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

58. (New) The method of claim 31 wherein said compound is 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

59. (New) The method of claim 31 wherein said compound is 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

60. (New) The method of claim 31 wherein said compound is 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

61. (Cancelled)

62. (New) The method of claim 35 wherein said compound is 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole.

63. (New) The method of claim 35 wherein said compound is 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

64. (New) The method of claim 35 wherein said compound is 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

65. (New) The method of claim 35 wherein said compound is 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole.

66. (New) The method of claim 35 wherein said compound is 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

67. (New) The method of claim 35 wherein said compound is 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

68. (New) The method of claim 35 wherein said compound is 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

69. (New) The method of claim 35 wherein said compound is 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

70. (New) The method of claim 35 wherein said compound is 4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole.

71. (First Amendment) The method of claim 40 wherein said compound is 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole.

72. (First Amendment) The method of claim 40 wherein said compound is 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

73. (First Amendment) The method of claim 40 wherein said compound is 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

74. (First Amendment) The method of claim 40 wherein said compound is 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole.

75. (First Amendment) The method of claim 40 wherein said compound is 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

76. (First Amendment) The method of claim 40 wherein said compound is 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

77. (First Amendment) The method of claim 40 wherein said compound is 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl) pyrrole.

78. (First Amendment) The method of claim 40 wherein said compound is 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole.

79. (Cancelled)

80. (Cancelled)

81. (Cancelled)

82. (Cancelled)

83. (Cancelled)

84. (Cancelled)

85. (Cancelled)

86. (Cancelled)